K021941

JUL - 2 2002

XIII. Special 510k Summary

Contact Person: Bill Mobbs, Managing Director

Date of Summary Preparation: June 5, 2002

Trade Name: ITL SampLok® Luer Kit

Common Name: Needle holder and luer adaptor for blood sample collection for testing

Classification Name: Tubes, vials, systems, serum separators, blood collection

Establishment Registration Number: 9033436

Class: II

Product Code: JKA

Legally Marketed Substantially Equivalent Devices: SampLok® Needle Holder (K000777) and Nipro Luer Adaptor (K992729)

Description of Device: Luer Kit for use with blood collection systems for collecting blood samples ~~from Y connectors.~~

Intended Use: The ITL SampLok® Luer Kit is intended to be used as part of a vacuum blood collection equipment for the collection of blood samples for various types of blood tests.

Comparison of Technical Characteristics: The ITL SampLok® Luer Kit consists of preattached SampLok® Needle Holder and Nipro Luer Adaptor, individually packaged and sterilized using ETO. The individual components are identical to previously cleared devices. The difference is that the previously cleared devices were packaged separately. The ITL SampLok® Luer Kit packages these two accessories together.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JUL - 2 2002

Mr. William Mobbs
Director
ITL Corporation
41-45 Tennant St, Fyshwick,
Canberra, ACT 2609
AUSTRALIA

Re: K021941
Trade/Device Name: ITL SampLok® Luer Kit
Regulation Number: 872.1675 and 880.5570
Regulation Name: Blood Specimen Collection Device and Hypodermic Single Lumen Needle
Regulatory Class: II
Product Code: JKA and FMI
Dated: June 6, 2002
Received: June 13, 2002

Dear Mr. Mobbs:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4618 . Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): K021941

Device Name:_____

Special 510(k) Device Modification	ITL Corporation Pty Ltd
ITL SampLok® Luer Kit	6/7/02

XII. Indications for Use

The ITL SampLok® Luer Kit is intended to be used as a part of vacuum blood collection equipment for the collection of blood samples for various types of blood tests..

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Rebecca Cicconetti

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K021941

Prescription Use ✓ OR **Over-The-Counter Use**___
(Per 21 CFR 801.109)

(Optional Format 1-2-96)